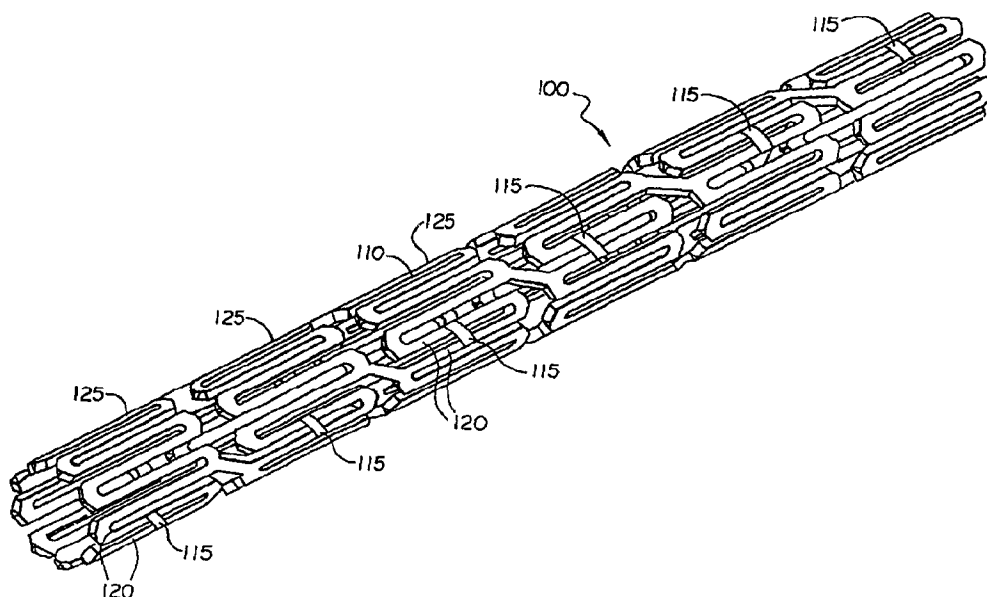




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : A61F 2/06	A1	(11) International Publication Number: WO 00/41649 (43) International Publication Date: 20 July 2000 (20.07.00)
(21) International Application Number: PCT/US00/00603 (22) International Filing Date: 11 January 2000 (11.01.00) (30) Priority Data: 09/232,404 15 January 1999 (15.01.99) US (71) Applicant: SCIMED LIFE SYSTEMS, INC. [US/US]; One SciMed Place, Maple Grove, MN 55311-1566 (US). (72) Inventor: KOCUR, Gordon J.,; 508 Lonesome Pine Trail, Lino Lakes, MN 55014 (US). (74) Agent: GRAD, Jonathan; Vidas, Arrett & Steinkraus, 6109 Blue Circle Drive, Suite 2000, Minnetonka, MN 55343-9131 (US).		(81) Designated States: CA, JP, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>

(54) Title: STENTS WITH TEMPORARY RETAINING BANDS



(57) Abstract

A self-expanding stent is formed of a stent framework (110) and retaining segments (115) disposed about the framework. The retaining segments hold the stent in a partially or fully contracted state in a bodily vessel. The retaining segments are constructed to fail after a predetermined time or upon the application of a predetermined force. Upon failure of the segments, the stent expands in the bodily vessel.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

STENTS WITH TEMPORARY RETAINING BANDS

U.S. Application No. 09/232,404 from which this application claims priority, is incorporated in its entirety herein by reference.

5

BACKGROUND OF THE INVENTION

This invention relates specifically to stents but also generally to any expandable, implantable endoprosthesis such as grafts, stent-grafts and the like. When the term 'stent' is used herein, it is to be understood in a general sense as including all such expandable prostheses, unless otherwise indicated.

The use of endoprotheses such as stents, stent-grafts and grafts is well known in maintaining the patency of bodily vessels including blood vessels and biliary ducts. Typically, an endoprosthesis is implanted in a vessel which has been occluded, which is subject to an aneurysm, which has a lesion present or is otherwise damaged. Often, during the implantation of the endoprosthesis, the vessel will suffer from trauma. The trauma may be as a result of the dilation prior to the implantation of the endoprosthesis, the presence of a foreign body (the endoprosthesis) in the bodily vessel or as a result of other causes. Regardless of the source of the trauma, the vessel may be in a weakened and inflamed state as a result of implantation of the endoprosthesis. Although it is desirable to maintain the vessel at as large a diameter as possible to minimize the possibility of restenosis, the weakening of the vessel resulting from the trauma limits the extent to which the vessel can be dilated.

The endoprotheses that are currently available are typically balloon expandable, self-expanding, or balloon assisted self-expanding devices. Balloon expandable stents achieve their maximum diameter upon expansion by a balloon. Once they have been seated in the vessel, they are incapable of further expansion unless a balloon is reinserted in the stent and expanded. Self-expanding and balloon assisted expandable stents, on the other hand, continually exert an outward force as they try to attain their maximum possible diameter. Thus, even after the stent is implanted, if it has not reached its maximum diameter it continues to try to open further exerting a force on the vessel walls.

It would be desirable to provide an endoprosthesis which has some of the characteristics of balloon expandable stents following deployment and which has some

of the properties of self-expanding stents after a predetermined period of time or after the application of a predetermined amount of force thereto. In particular, it is desirable to provide an endoprosthesis which does not impart to the vessel walls the outward forces associated with a self-expanding stent while the vessel is recovering from the trauma of the deployment procedure and yet provides the outward expanding force of a self-expanding stent when the vessel is sufficiently recovered from the trauma.

It is also desirable to provide an endoprosthesis which only partially expands or does not expand at all during deployment and which expands at some later point following deployment.

The present invention provides such an endoprosthesis.

For the purposes of this disclosure, unless otherwise indicated, the term 'degradation' shall refer to degradation in its ordinary sense as well as biodegradation, erosion, and dissolution.

BRIEF SUMMARY OF THE INVENTION

The invention is directed to an expandable stent comprising an expandable stent framework which is expandable from a reduced diameter configuration to a fully expanded configuration. At least one stent retaining segment which is constructed to fail is disposed about the stent framework to maintain the stent framework in a reduced diameter state until the retaining segment fails. The retaining segment may be constructed to fail after a predetermined period of time has elapsed and/or after a predetermined amount of force has been applied to the stent. The retaining segment may be entirely biodegradable, biodegradable in part, erodible or made of an inert material with fatigue points.

In one embodiment of the invention, the retaining segment forms a band around the outside of the stent or interwoven about the stent. In particular, the retaining segment may be woven such that it alternates on the inner and outer strut surfaces.

The retaining segment may also be provided in the form of a web.

The invention is also directed to treatment methods in which the inventive stent is deployed and further expands upon failure of the retaining segments. In the case of an inert segment, failure of the retaining segment may result from mechanical failure. Stresses on the stent which are likely to result in or contribute to the desired failure include pulsing blood pressure which is thought to cause an alternating

lengthening and shortening of the stent.

In the case of a biodegradable segment or an erodible segment, failure results from degradation or erosion of the segment. Finally, in the case of a segment made of inert material and connecting dots of biodegradable material, failure results
5 from degradation or erosion of the connecting dots.

More generally, the invention is directed to the application of a secondary increased expansive force to a body lumen by an implantable expandable medical device such as a stent endoprostheses including stents, stent-grafts, grafts and vena cava filters at a predetermined time following implantation into a body lumen.

10

BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWING(S)

Figure 1 shows a perspective view of an inventive stent.

Figure 2 shows a perspective view of another inventive stent.

Figure 3 shows a perspective view of an inventive stent illustrating
15 several types of retaining segments.

Figure 4a shows a perspective view of another inventive stent.

Figure 4b shows a perspective view of another inventive stent.

Figure 4c shows a cross-sectional view of the stent of Fig. 4b along the
lines 4c-4c.

20 Figures 5a-c show retaining segments with fatigue points.

Figure 6a shows a perspective view of a retaining segment.

Figure 6b shows a perspective view of another retaining segment.

Figures 7 shows a perspective view of another embodiment of the
inventive stent.

25 Figure 8 shows a perspective view of another embodiment of the
inventive stent.

Figure 9 shows a partial cross-section of a stent with a retaining segment fused to the stent.

Figure 10 shows a partial cross-section of a stent with a retaining
30 segment bonded the stent.

Figures 11a-e are schematic representations of stent with retaining segments in the form of webs.

Figure 12 is a perspective view of a die for use in the present invention.

Figures 13a-d are schematic representations of a stent with a retaining band prior to (Figs. 13a and 13c) and following (Figs. 13b and 13d) failure of the retaining segment.

5 DETAILED DESCRIPTION OF THE INVENTION

While this invention may be embodied in many different forms, there are described in detail herein specific preferred embodiments of the invention. This description is an exemplification of the principles of the invention and is not intended to limit the invention to the particular embodiments illustrated.

10 With reference to the Figures in general and Fig. 1 in particular, the present invention is directed to an expandable stent, shown generally at 100, comprising an expandable stent framework 110, and a stent retaining segments 115 which are selected portions of the stent framework. Expandable stent framework 110 is a self-expanding stent which is expandable from a reduced diameter configuration to a fully
15 expanded configuration. Stent retaining segments 115 maintains the stent framework in a less than fully expanded configuration until it fails. In the embodiment of Fig. 1, one pair of struts 120 per stent section 125 are joined together with a retaining segment 115. Each stent retaining segment 115 is circumferentially offset from the adjacent stent retaining segment by one strut 120.

20 The invention also contemplates providing a plurality of stent retaining segments per stent section. Figure 2 shows stent 100 with two stent retaining segments 115 per stent section 125. Stent retaining segments 115 are oppositely disposed from one another across stent section 125.

Figure 3 illustrates several other configurations of stent retaining
25 segments. In stent section 125a, stent retaining segments 115a join every other pair of adjacent struts 120 in the center of the struts. In stent section 125b, every pair of adjacent struts are joined by stent retaining segments 115b. Stent retaining segments 115b are seen to alternate between a first end of adjacent and a second end of adjacent struts. The stent of Figure 3 also shows a retaining segment 115d connecting adjacent
30 stent sections 125d and 125e. The stent of Figure 3 also shows a retaining segment 115e extending between three adjacent struts. It is understood that the stent of Figure 3 is intended to be illustrative of several embodiments. Those of ordinary skill in the art will recognize that the invention contemplates stents in which a desired number of stent

sections have stent retaining segments as in section 125a. Similarly, the invention contemplates stents having stent retaining segments arranged entirely as in section 125b. It will also be recognized that the invention contemplates stents with interconnected stent sections as in sections 125d and 125e. Finally, it will be recognized that each
5 retaining segment may restrain multiple struts beyond the two shown in the figure.

The use of small retaining segments is beneficial for several reasons. First, retaining individual segments of the stent allows for greater stent flexibility. Second, using relatively small retaining segments results in a reduced negative tissue response to the implant and the segment in particular. Third, retaining only stent
10 segments allows for better control over stent expansion and thus, how the additional secondary force is applied to the body lumen.

The invention also contemplates the use of larger stent retaining segments in the form of stent retaining bands. In one embodiment of the invention, the stent retaining segments form one or more bands extending around the circumference.
15 The stent of Fig. 4a, shown generally at 100, comprises an expandable stent framework 110, and a stent retaining segment 115 which is disposed about the circumference of the stent 110 in the form of a band. Expandable stent framework 110 is a self-expanding stent which is expandable from a reduced diameter configuration to a fully expanded configuration. Stent retaining segment 115 maintains the stent framework in a less than
20 fully expanded configuration until it fails. The invention also contemplates the possibility of providing the stent with additional retaining bands beyond the two shown in Fig. 4a.

The band-like segments may extend around the stent from the outside of the stent or may be interwoven around the circumference of the stent as shown in Figs.
25 4b and 4c. Bands 115 weave in-between struts 120.

The invention contemplates several different types of stent retaining segments. One stent retaining segment suitable for use in the present invention is shown in Fig. 5a. Stent retaining segment 215 has a single fatigue point 220. Fatigue point 220 is a narrowed region of the band. Fig. 5b shows a stent retaining band 215 with two
30 fatigue points 220. The shape of the fatigue points shown in Figs. 5a and 5b is intended to be for illustrative purpose only. Fatigue points of other geometries and configurations may also be used. A retaining band with a single fatigue point may also be used as may a retaining band with additional fatigue points. Fig. 5c shows a retaining

segment 215 with two holes 223 punched therethrough. Fewer or additional holes may also be present in the region to weaken the segment.

Retaining band 215 is made of an inert or biostable material such as polytetrafluoroethylene (PTFE). Other inert or biostable materials may also be used.

- 5 Desirably, a material with a low tissue response such as PTFE, polyurethanes, polyesters, and silicones will be used.

Another suitable stent retaining segment is shown generally at 315 in Fig. 6a. Retaining segment 315 is formed of two segments of inert or biostable material 325 and a segment of biodegradable material 320 joined together. Such a design may also
10 be used for a stent retaining band as shown in Fig. 6b. Retaining band 315 is formed of a segment of inert or biostable material 325 and a segment of biodegradable material 320 joined together. As shown in Figs. 6a and 6b, biodegradable segment 320 is provided in the form of a dot. Biodegradable segment 320 may suitably be provided in other shapes as well. In particular, the retaining segment or band may be provided with
15 the same general shape as bands 215 shown in Figs. 5a and 5b, with the fatigue point replaced by a biodegradable segment where the band narrows. An embodiment of retaining band 315 having two inert segments 325 and two biodegradable segments 320 is shown in Fig. 6c. A retaining segment may be similarly prepared with multiple biodegradable segments. The invention further contemplates the use of additional inert
20 segments and additional biodegradable segments.

Another embodiment of the invention is shown in Fig. 7. Stent 100 comprises an expandable stent framework 110, and a stent retaining band 115 which is disposed about the stent the circumference of the stent 110. Expandable stent framework 110 is a self-expanding stent which can expand from a reduced diameter
25 configuration to a fully expanded configuration. Stent retaining band 115 is disposed about stent framework 110 at an oblique angle relative to the longitudinal axis 130 of the stent. The invention is not intended to be limited to the specific oblique angle shown but rather, is intended to encompass the full range of oblique angles. The stent may have fewer or more retaining bands than shown in Fig. 7.

30 Another embodiment is shown in Fig. 8. Stent 100 is retained in a reduced diameter configuration via stent retaining band 115 which is helically disposed about the stent.

The invention also contemplates the use of retaining structures such as webs or nets to maintain the stent in a reduced configuration or less than fully expanded configuration. In general the web is of a woven or closed cell design that is fused to the exterior of the stent. The web can be in the shape of a tube that wraps around individual
5 segments of the stent as shown generally at 415 in Fig. 11a. The web can also wraps around the entire length of the stent as shown in Fig. 11b. Web 415 is disposed about the length of stent framework 110. The web desirably has failure points 420 to allow the web to pull apart following failure.

The web may also be a flat sheet as shown at 415 in Fig. 11c. Web 415
10 covers a group of cells like a patch. The web patch may either have failure points such as fatigue points or degradable portions to release the stent. The web patch may be attached to the stent struts via a degradable material as shown in Fig. 11d. Web 415 is attached to stent 110 via degradable adhesive 122.

Another web configuration is shown at 415 in Fig. 11e. Web 415 is in a
15 hexagonal pattern with fatigue points 420 at the vertices of the hexagons. Other polygonal configuration may be substituted.

As with the retaining segments, the web may be made of a biodegradable material or may be made of an inert material with fatigue points. An inert material with biodegradable connecting material may also be used.

20 Although specific web configurations have been shown herein, those of ordinary skill in the art will understand these to be exemplary. Other web configurations may be used as well.

Retaining segments in the form of thin strips may be cut from sheets of a desired material. Threads of retaining segment may also be made by extrusion. Fatigue
25 points may be made by varying the extrusion rate of the thread or by varying the tension on the thread during extrusion. Tension may also be used to introduce fatigue points into other types of retaining segments such as band-shaped retaining segments. Materials such as PTFE will neck under tension thereby forming a fatigue point. Fatigue points may also be placed in the material by cutting the material with a laser or
30 with a punch as illustrated in Fig. 5c. Other methods for introducing fatigue points in a retaining segment include the spot application of a laser, of heat, of one or more solvents or other chemicals such as etchants, or of radiation to weaken or make the

material brittle. Fatigue points may also be introduced into a retaining segment via the use of a second material in the segment.

As shown in some of the figures, the retaining segment is disposed on the outside of the stent. *In situ*, the retaining band is disposed between the vessel wall and the stent thereby facilitating encapsulation of the retaining band in tissue and preventing the inert material from being carried away in the vessel. This is particularly desirable where inert material is used for the retaining band.

The retaining segment may be secured to the stent in a variety of ways. Thread-like and other suitably sized retaining segments may be woven through the stent and/or tied to the stent or struts of the stent.

Alternatively, as illustrated in Fig. 9, a piece of retaining segment 115a may be placed on an outer surface of a strut 120a and another piece of retaining segment 115b may be placed on an inner surface of a strut 120b. The two pieces may then be fused together at 121 by heat, solvent, or adhesive.

Other methods of attachment include looping a retaining segment over a strut and securing the segment to itself or the strut melting with heat or solvent or adhesive. The retaining segment may also be wrapped around struts or layered around struts or attached directly to struts and secured thereto by applying heat or a solvent or other suitable adhesives including bioabsorbable, biodegradable or erodible compositions such as those disclosed below. As shown in Figs. 4c and 10, a biostable material 122 may also be spot melted to or around a strut 120 and then fused to a retaining segment 115 made of biodegradable material by heat, solvent or adhesive. Where a solvent is used, excess solvent may be evaporated off or otherwise driven off.

One method of attachment involves providing a mandrel with degradable dots of material thereon. A stent is then disposed about the mandrel and a PTFE strip placed over a portion of the stent. Desired portions of the stent may be clipped so as to retain the stent in a reduced configuration on the mandrel. Heat is applied to the stent so as to fuse the dots to the stent and the PTFE. The stent is then cooled and removed from the mandrel.

In another method for applying a retaining segment to a stent, retaining segments are placed in a die which is then disposed about a stent in a reduced diameter configuration. The stent is then expanded. Specifically, a circular die at a desired intermediate stent diameter is provided. The die is provided in two halves, one of which

is die, shown at 500 in Fig. 12, with groves 501 cut into the inner diameter surface for the retaining segment pattern. Retaining segments of a desired material are placed into the groves in the die halves and the die halves reassembled. A mandrel is provided and a nitinol stent is reduced in size onto the mandrel by cooling the stent below the austenitic-martensitic transition temperature to make it martensitic. Optionally, the mandrel may have grooves cut therein in the retaining segment pattern for placement of a desired material on inner diameter surface of stent. The reduced stent and mandrel are placed into the die. The stent is then heated to expand it inside the die. Adhesives, heat or solvent are then used to fuse the retaining segments to each other and/or the stent struts, retaining the stent at the desired diameter.

In yet another method of applying a retaining segment to a stent, clips may be used to hold struts or cells in the closed position in one or more desired portions of the stent. Threads of retaining segment material are laid across these portions of the stent and attached to the struts using adhesives, heat, or solvent.

Retaining segments may also be secured to a stent by hot extruding strands of material between confined struts (i.e. struts in a closed position or reduced diameter position).

Retaining segments may also be applied by first holding the stent in a desired reduced diameter configuration via the use of a retaining die or clamps. Strands of the retaining material may then be sprayed on the interior of the stent or the exterior of the stent in the area to be retained.

Figures 13a-13d illustrate a portion of a stent with a retaining segment prior to and after failure. As shown in Figs. 13a and 13c, stent 110 has a retaining segment 115 which retains at least a portion of the stent in a reduced configuration. Segment 115 is attached to the stent at several points 122. Upon failure of the retaining segment 115, as shown in Figs. 13b and 13d, stent 110 expands.

Suitable biodegradable or bioabsorbable materials for use in the retaining segments, bands or webs include:

Poly(L-lactide) (PLLA), Poly(D,L-lactide) (PLA), poly(glycolide) (PGA), poly(L-lactide-co-D,L-lactide) (PLLA/PLA), poly(L-lactide-co-glycolide) (PLLA/PGA), poly(D,L-lactide-co-glycolide) (PLA/PGA), poly(glycolide-co-trimethylene carbonate) (PGA/PTMC), polydioxanone (PDS), Polycaprolactone (PCL), polyhydroxybutyrate (PHBT), poly(phosphazene) poly(D,L-lactide-co-

caprolactone) (PLA/PCL), poly(glycolide-co-caprolactone) (PGA/PCL), poly(phosphate ester), poly(DL-lactic acid), poly(glycolic acid).

PGA and PLLA/PGA are particularly desirable materials. Other suitable materials include collagen, fibrin, and cellulose.

5 More generally, general classes of degradable material include polyesters, polyamides, polyanhydrides and polyorthoesters. The latter two are surface erodible types. All of these are exemplary only.

As discussed above, the retaining segments may be designed to last for a predetermined period of time prior to failing. Where the retaining segment incorporates
10 biodegradable materials, the lifetime of the segment may be altered by altering the chemical structure of the biodegradable portion of the segment. Hydrolysis is the basic reaction for the most biodegradable polymers. The hydrolytic degradation rate can be altered several thousand-fold by changing the chemical structure in the polymer backbone. For example, aliphatic polyanhydrides degrade in a few days while the
15 aromatic polyanhydrides degrade over a period of a few years. Polyorthoester is a slow surface eroding material. In the presence of acid additive, a so-called excipient, it has a faster degradation rate. In contrast, in the presence of basic substance, it suppresses degradation. When longer lasting retaining segments are desired, the aromatic polyanhydride and non-additive polyorthoester will be preferred segment materials.

20 Also, for example, polymers that contain hydrolytically labile linkages in their backbone can hydrolyze by two different mechanisms. These are bulk erosion and surface erosion. In a bulk eroding polymer, the hydrolytic process occurs throughout the matrix of the polymer whereas in surface erosion the hydrolysis is only confined to the outer layer of the polymer. Thus, the latter is especially preferred when longer
25 degradation is desired.

Specific segment materials include, for example, poly(ortho esters) such as 50:50 HD/t-CDM (1,6-Hexanediol-co-trans-Cyclohexanedimethanol) poly(ortho ester) with 0.2% poly(sebacic anhydride) excipient. Polyanhydrides such as poly[bis(p-carboxyphenoxy)propane anhydride] (PCPP) and poly(terephthalic anhydride) (PTA)
30 may also be used.

If a polyanhydride is selected as the segment material, for example, its thickness can be selected such as to control degradation time. Thus, a segment may be tailored to have a degradation time of a week, a month, two months, or up to six months,

or any other suitable period of time. Segment material of PGA, for example, may be provided at a selected thickness to provide a degradation period of about two weeks, for example.

The retaining segment is desirably designed to fail after a predetermined period of time following implantation in the bodily lumen. Failure includes mechanical failure and failure resulting from the biodegradation of at least a portion of the segment. Retaining segments which fail after about one week in the lumen are desirable as are retention segments that fail after about one month in the lumen. More desirably, the retention segments will last from about one to about six months following implantation. This allows the safe application of additional force to the body lumen after the initial tissue response to the implant and lumen remodeling has progressed or at a time where additional external force on the body lumen is expected to increase as in the case of an adjacent growing tumor.

The retaining segments may also be designed to fail after the application of a predetermined amount of force such as by a balloon.

The invention also contemplates providing the retaining segment with a treatment or therapeutic agent. A retaining segment material that elutes the therapeutic agent can be made by dissolving or suspending the therapeutic agent(s) and retaining segment polymer in a solvent. Threads may then be extruded and films may be sprayed or cast for making strips or webs for use as retaining segments. The treatment agent may also be provided by any other suitable techniques including impregnating or coating the segment with the treatment agent.

Desirable types of therapeutic agents include anti-inflammatory agents, anti-proliferative agents, anti-platelet agents, anti-thrombin agents, anti-oxidant agents, gene therapy agents and suitable combinations of the above agents.

Suitable treatment agents include drugs such as radiochemicals to irradiate and prohibit tissue growth and human growth factors including VEGF (Vascular Endothelial Growth Factor), TGF-beta (Transforming Growth Factor-beta), IGF (Insulin-like Growth Factor), PDGF (Platelet-derived Growth Factor) and FGF (Fibroblast Growth Factor), etc. The drug can be an anticoagulant, e.g. D-Phe-ProArg chloromethyl ketone. An RGD (Arginine-Glycine-Aspartic Acid) peptide-containing compound, heparin, antithrombin compounds, platelet receptor antagonists, antibodies, aspirin, prostaglandin inhibitors, platelet inhibitors, or antiplatelet peptide may also be used.

The drug can be an inhibitor of vascular cell growth, DNA, RNA, cholesterol-lowering agents, vasodilating agents. Other suitable drugs include Taxol™ (paclitaxel), 5-Fluorouracil, Beta-Estradiol. Metabolites and other derivatives of the above drugs or compounds may also be used. Any treatment agent may be used, singly or in
5 combination.

The treatment agent may be impregnated into the retaining segments or may be provided as a coating on the segments. The treatment agent may be applied by immersing the segments in the treatment agent or spraying the agent onto the segments. Suitable techniques for applying the treatment agent may be found in US application
10 08/874190 incorporated herein in its entirety by reference.

The invention is also directed to an expandable stent for implantation in a bodily lumen comprising an expandable stent framework and a stent retaining segment which is constructed to fail. As discussed above, the retaining segment may be designed to fail after a predetermined period of time such as about one week or about one month.
15 The retaining segment may also be designed to fail upon the application thereto of a predetermined pressure such as that provided by a balloon.

The inventive stents, grafts and stent-grafts may be formed from any suitable self-expanding and balloon assisted self-expanding stent or graft known in the art. One such suitable self-expanding stent is disclosed in commonly assigned US
20 Application No. 08/511076 filed August 3, 1995, incorporated herein in its entirety by reference. Another suitable stent is the Wallstent stent, described in US 4,655,771, US 4,954,126 and US 5,061,275 incorporated herein in their entirety by reference.

Suitable grafts may be formed by applying a graft material such as polytetrafluoroethylene (PTFE) or polyethylene terephthalate (PET) to the stent and
25 securing it thereto by an appropriate means such as sutures. Other graft materials including polyurethane, collagen, silicone, polypropylene and polyolefin may also be used. The graft material may be used as inner liner for the stent or as an outer liner for the stent. Other types of self-expanding grafts may also be used in the practice of the invention.

30 The invention is intended to be used with self-expanding endoprostheses in general and as such is not limited to coronary endoprostheses. Non-coronary applications include biliary prostheses and ureteral prostheses. In particular, the inventive endoprostheses may prove useful where it is necessary to maintain the patency

of vessels or ducts that are being closed by tumors including the bile duct and pancreatic tumors.

The invention provides endoprotheses which may partially expand or not expand at all during deployment and yet expand at a later point as a result of failure
5 of a retaining segment.

In another aspect, the invention is directed to a treatment method which comprises the steps of implanting an expandable stent with retaining segments such as those disclosed above, in a bodily lumen. The retaining segments may be constructed to fail by biodegradation or erosion or mechanical failure after a predetermined period of
10 time in the body such as about week or about a month. In accordance with this treatment method, the stent is implanted in a less than fully expanded state and expands upon the passage of a predetermined period of time such as a week, a month or longer with the failure or dissolution of the retaining segments. The stent may be delivered to the desired bodily location using a stent delivery catheter such as those disclosed in US
15 5,534,007 and US 5,571,135 incorporated herein in their entirety by reference or any other suitable stent delivery catheter.

Desirably, the retaining segments will be constructed to fail after the vessel has sufficiently healed to be able to safely withstand the increment in force following the failure of the segments.

The inventive treatment method also contemplates the deployment of a
20 stent with retaining segments as disclosed above in a bodily lumen and the subsequent application of a predetermined force to the stent to more fully expand the stent. Additional force may be applied immediately after deployment of the stent as a 'touch-up' or after some suitable period of time following deployment (i.e. days, weeks or
25 months). The additional force necessary to break the retention segments may be applied via an inflatable balloon on a catheter.

The invention is also directed more generally to an expandable medical device which is held in a contracted or partially contracted state by one or more retaining structures such as the segments described above. Suitable medical device
30 include stents, stent-grafts, grafts and vena cava filters.

The above disclosure is intended to be illustrative and not exhaustive. This description will suggest many variations and alternatives to one of ordinary skill in this art. All these alternatives and variations are intended to be included within the

scope of the attached claims. In particular, the invention contemplates stents with multiple types of retaining segments (e.g. bands and segments linking only several struts together). As such, stents which are combinations of those shown in the Figures are intended. Those familiar with the art may recognize other equivalents to the specific
5 embodiments described herein which equivalents are also intended to be encompassed by the claims attached hereto.

CLAIMS:

1. An expandable stent comprising:
an expandable stent framework, the stent framework expandable from a reduced diameter configuration to a fully expanded configuration;
5 and at least one stent retaining segment disposed about the stent, the stent retaining segment maintaining the stent framework in a less than fully expanded configuration, the stent retaining segment constructed and arranged to fail upon degradation of at least a portion of the segment.
2. The stent of claim 1 comprising a plurality of stent retaining segments.
- 10 3. The stent of claim 1 wherein the retaining segment is formed of at least one segment of inert material and at least one segment of biodegradable material joined together.
4. The stent of claim 3 wherein the retaining segment comprises a plurality of alternating inert segments and biodegradable segments joined together.
- 15 5. The stent of claim 1 wherein the retaining segment is made of a biodegradable material.
6. The stent of claim 5 wherein the biodegradable material is provided with a treatment agent.
7. The stent of claim 1 wherein the retaining segment is provided with a treatment
20 agent.
8. The stent of claim 1 wherein the retaining segment is disposed on the outside of the stent framework.
9. The stent of claim 1 wherein the retaining segment is interwoven through the stent framework.
- 25 10. The stent of claim 1 wherein the retaining segment is constructed to fail after a predetermined period of time in the body.
11. The stent of claim 10 wherein the period of time is at least one month.
12. The stent of claim 10 wherein the retaining segment is in the form of a band disposed about the stent.
- 30 13. The stent of claim 1 wherein the stent framework comprises a plurality of interconnected struts and the retaining segment interconnects connects no more than two adjacent struts.
14. The stent of claim 1 wherein the retaining segment is in the form of a web.

15. An expandable stent comprising:
an expandable stent framework, the stent framework expandable from a reduced diameter configuration to a fully expanded configuration;
and at least one stent retaining segment disposed about the stent,
5 the stent retaining segment maintaining the stent framework in a less than fully expanded configuration, the stent retaining segment constructed and arranged to have at least one fatigue point thereon.
16. The stent of claim 15 the retaining segment having a plurality of fatigue points thereon.
- 10 17. The stent of claim 15 wherein the retaining segment narrows at the fatigue point.
18. The stent of claim 15 wherein the retaining segment is made of an inert, material.
19. The stent of claim 18 wherein the material is PTFE.
20. The stent of claim 15 wherein the stent retaining segment is constructed to fail
15 after a predetermined amount of time in a bodily lumen.
21. The stent of claim 15 wherein the retaining segment is constructed to fail after at least one month in a bodily lumen.
22. The stent of claim 15 where the stent retaining segment is constructed to fail upon the application thereto of a predetermined force.
- 20 23. A treatment method comprising the steps of:
i) implanting an expandable prosthesis in a bodily lumen, the expandable prosthesis comprising
1) an expandable prosthesis framework, the prosthesis
framework expandable from a reduced diameter configuration to a fully
25 expanded configuration; and
2) a prosthesis retaining segment which is constructed to fail
upon dissolution of at least a portion thereof, the segment formed at least in part of a biodegradable material
the prosthesis retaining segment disposed about the prosthesis
30 and maintaining the prosthesis framework in a reduced diameter configuration.
24. The method of claim 23 wherein the prosthesis retaining segment is constructed to fail after a predetermined time in a bodily lumen.

25. The method of claim 23 wherein the predetermined time is at least one month.

26. A treatment method comprising the steps of:

i) implanting an expandable prosthesis in a bodily lumen, the expandable prosthesis comprising

5 1) an expandable prosthesis framework, the prosthesis framework expandable from a reduced diameter configuration to a fully expanded configuration; and

 2) a prosthesis retaining segment which is constructed to have at least one fatigue point thereon to facilitate failure of the segment under a
10 predetermined set of conditions,

 the prosthesis retaining segment disposed about the prosthesis and maintaining the prosthesis framework in a reduced diameter.

27. An expandable medical endoprosthesis for implantation in a bodily vessel comprising:

15 an expandable prosthesis framework expandable from a reduced diameter configuration to a fully expanded configuration;

 and at least one endoprosthesis retaining structure attached to the endoprosthesis framework and disposed about the endoprosthesis framework, the endoprosthesis retaining structure maintaining the endoprosthesis framework in a
20 reduced diameter configuration.

28. The endoprosthesis of claim 27 wherein the retaining structure is in the form of a segment.

29. The endoprosthesis of claim 27 wherein the retaining structure is in the form of a web.

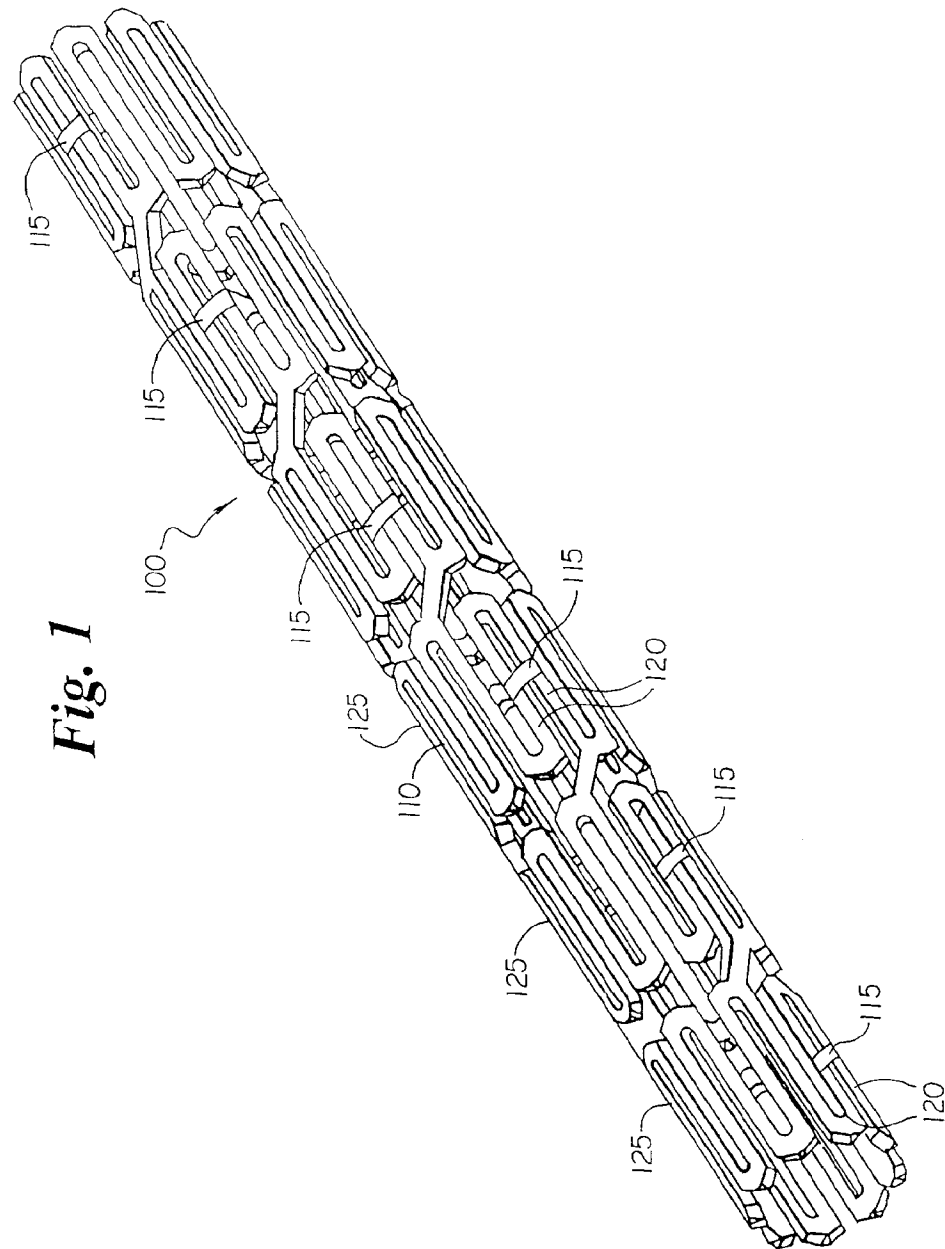


Fig. 1

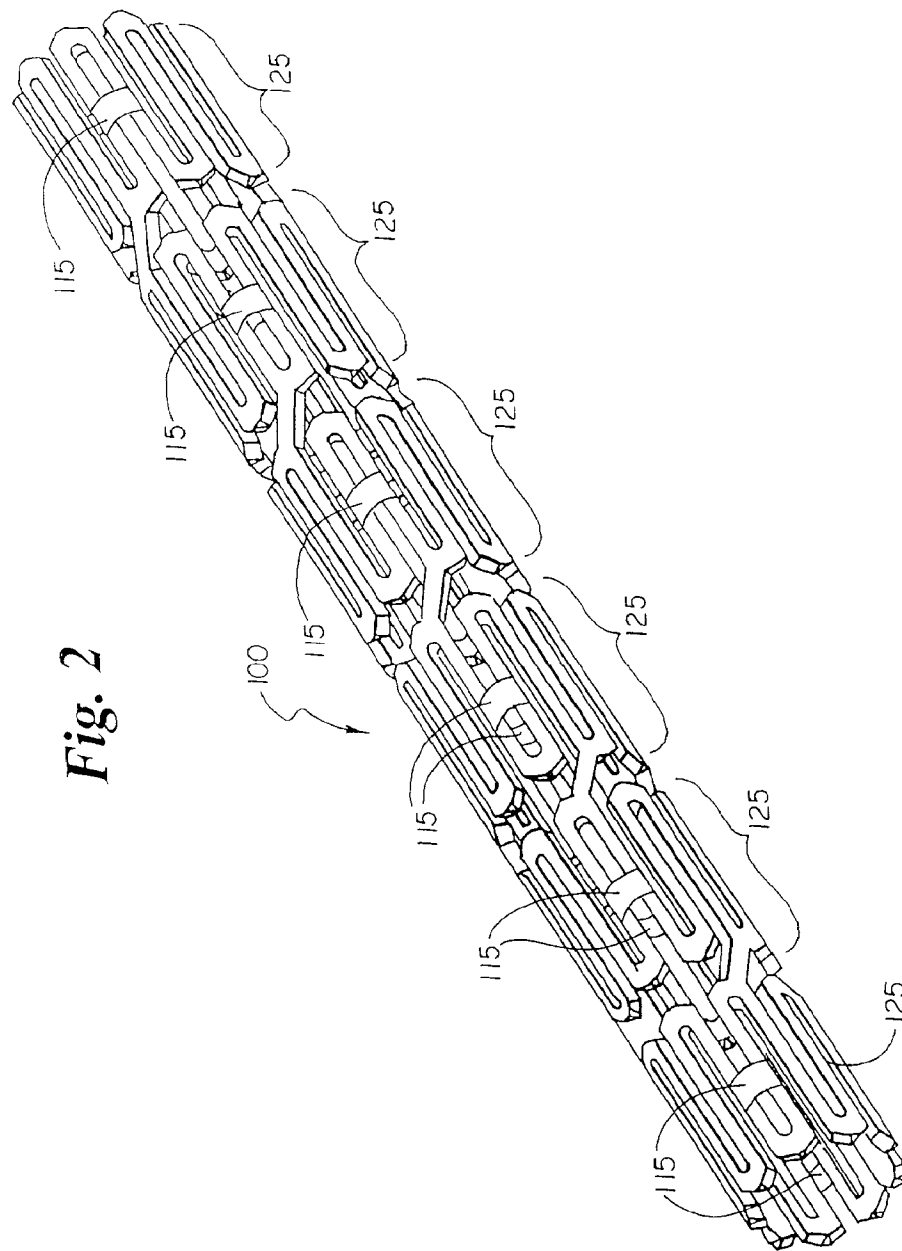


Fig. 2

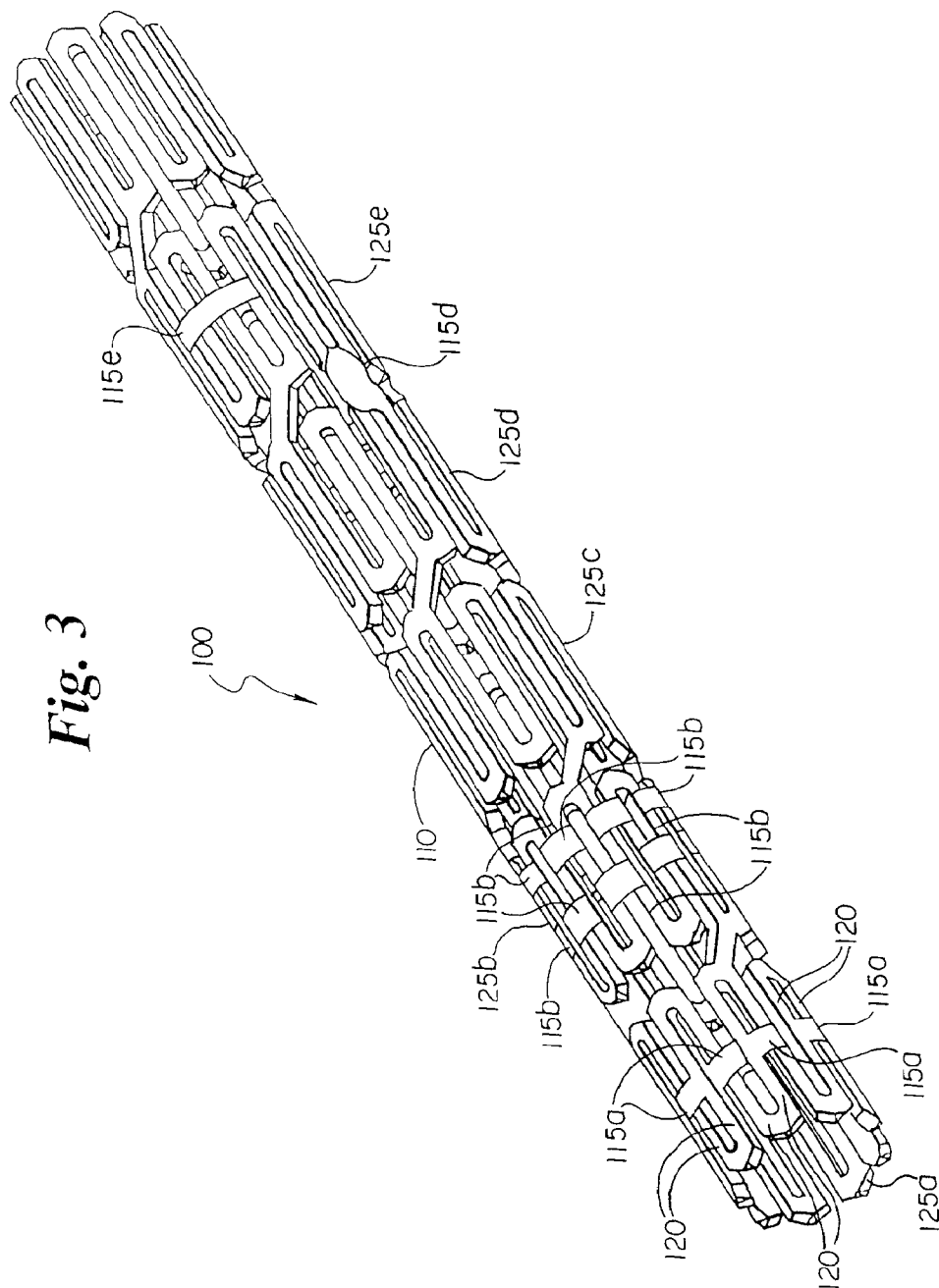
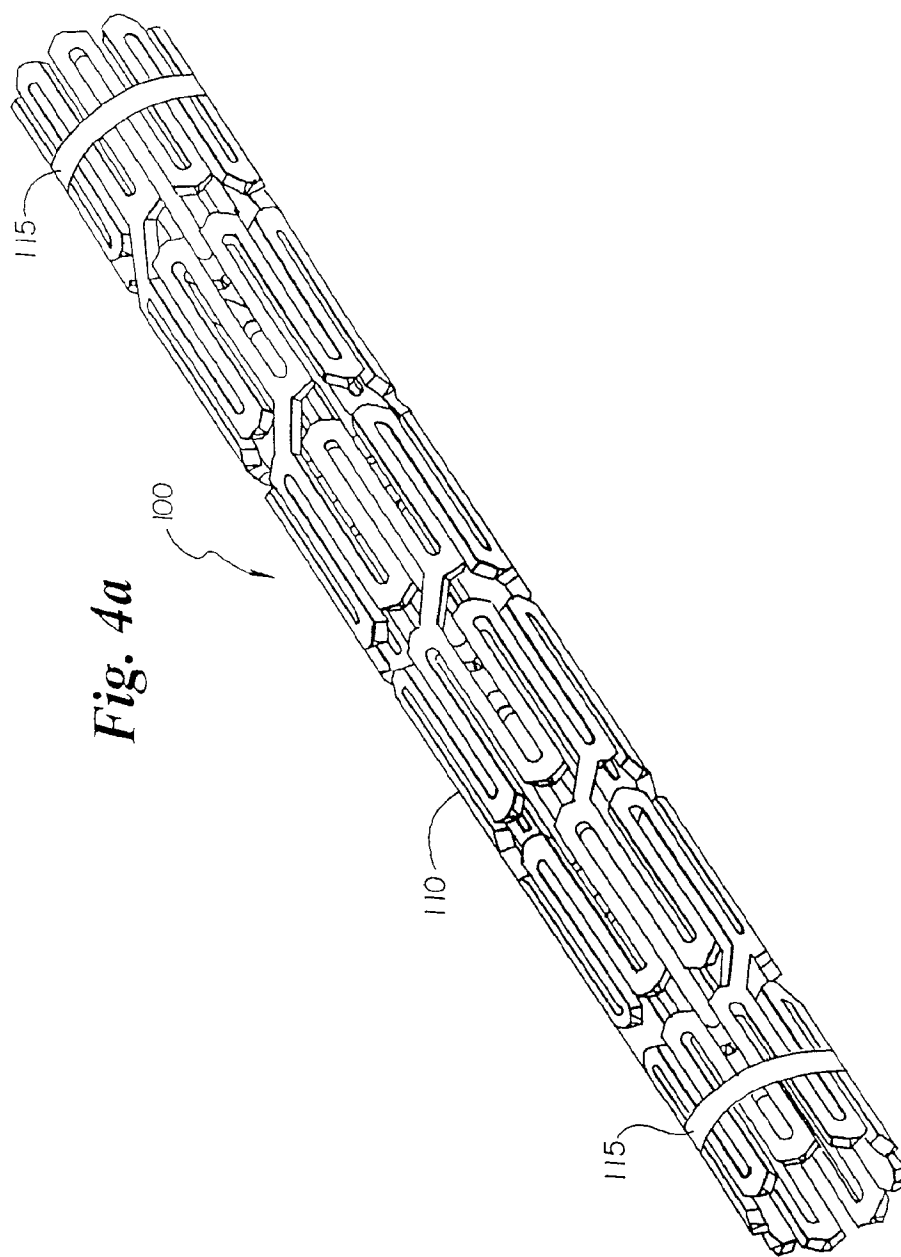


Fig. 3



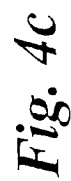
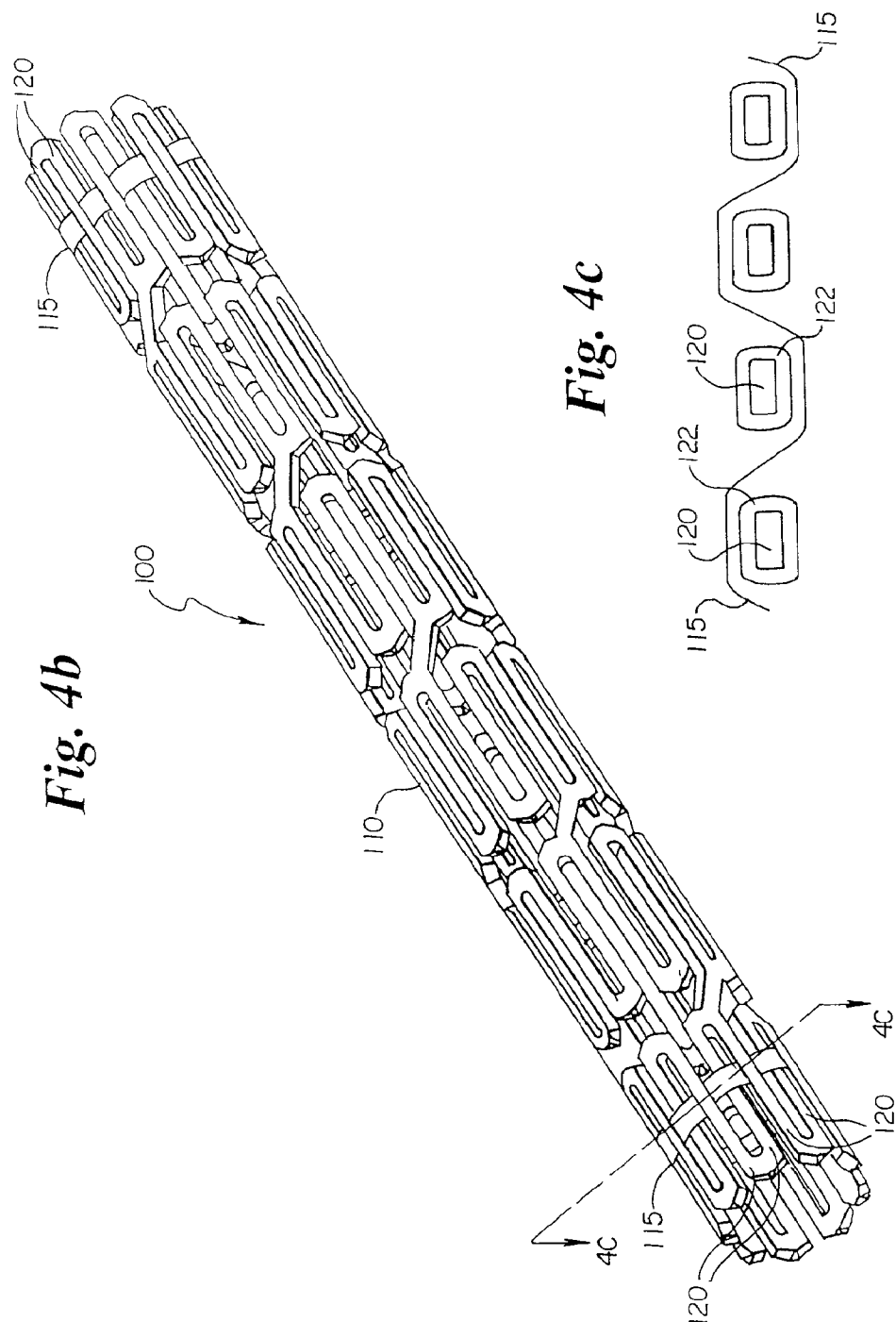


Fig. 5a

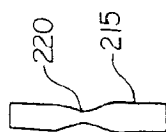


Fig. 5b

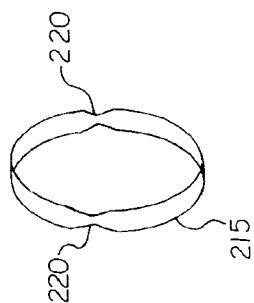


Fig. 5c

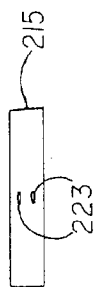


Fig. 6a

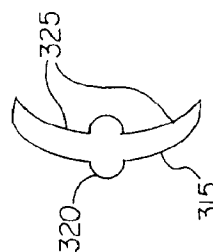


Fig. 6b

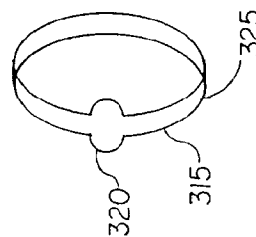
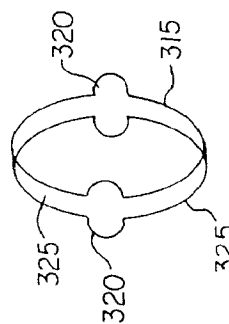
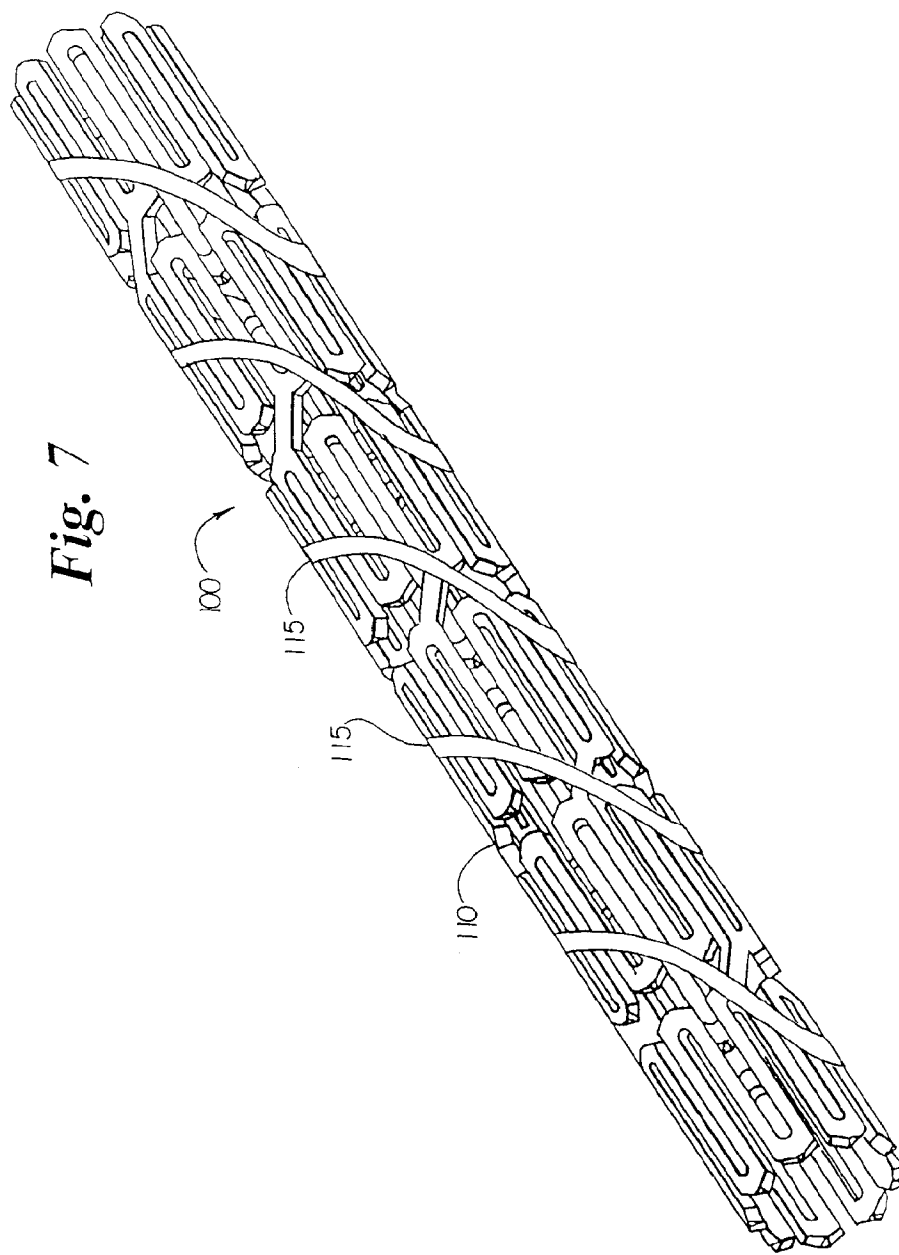


Fig. 6c





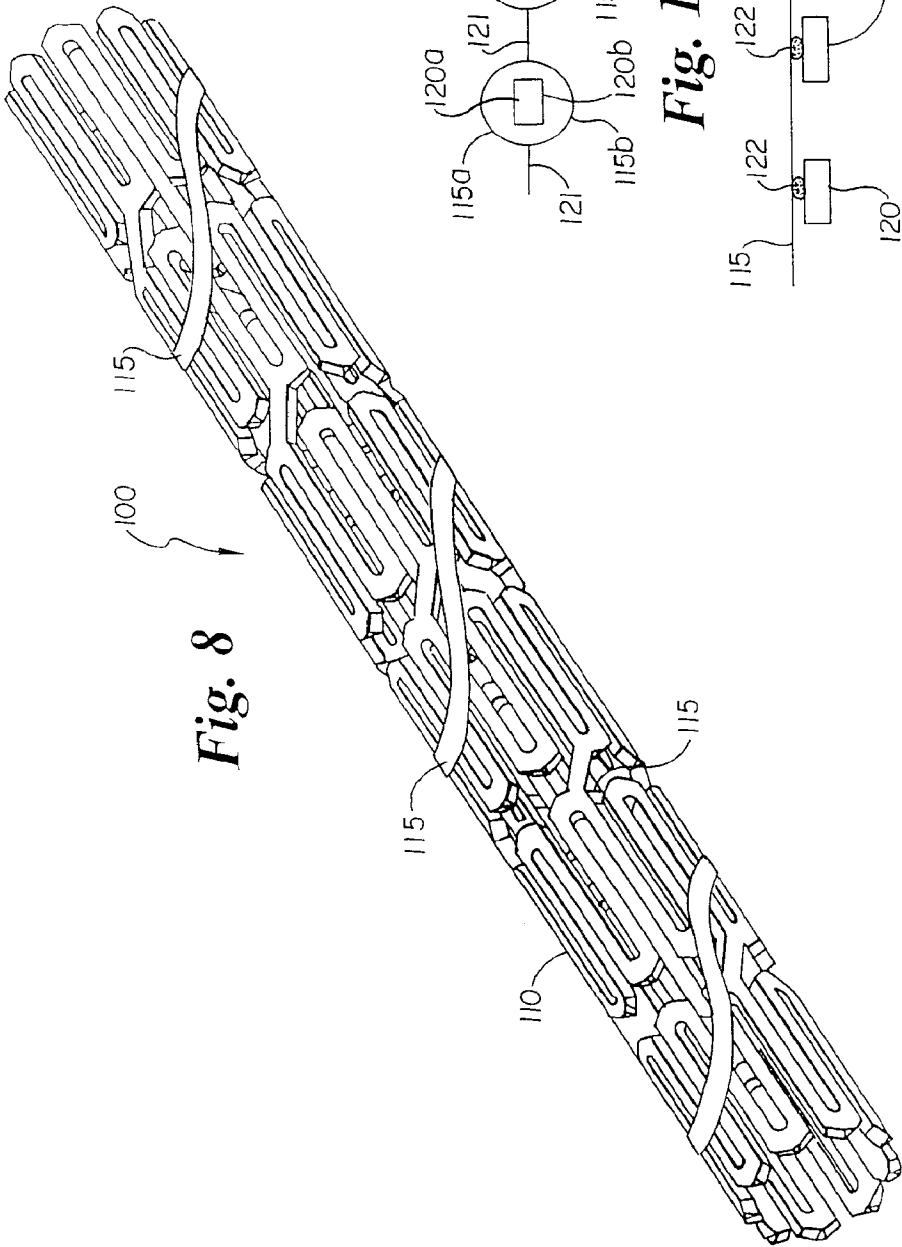


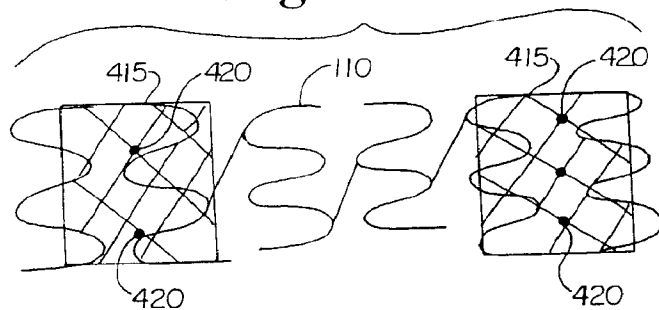
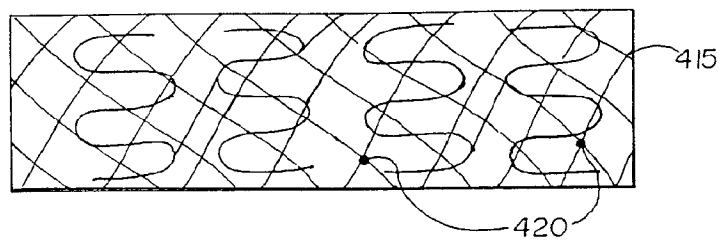
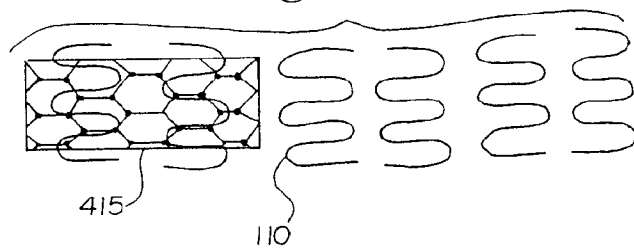
Fig. 11a*Fig. 11b**Fig. 11c*

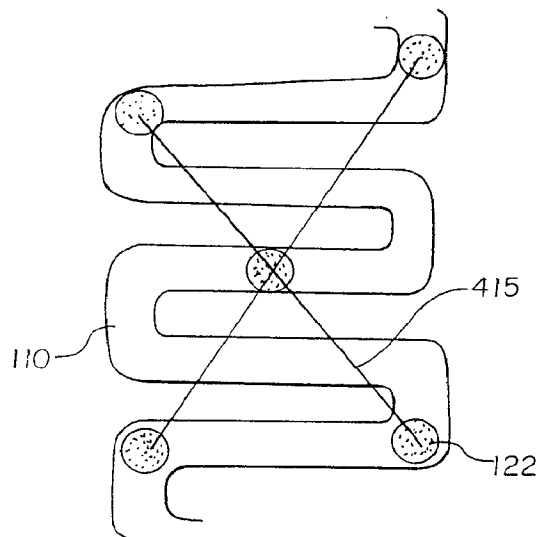
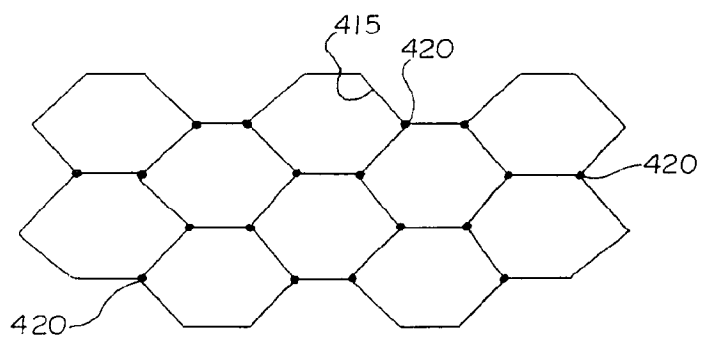
Fig. 11d*Fig. 11e*

Fig. 12

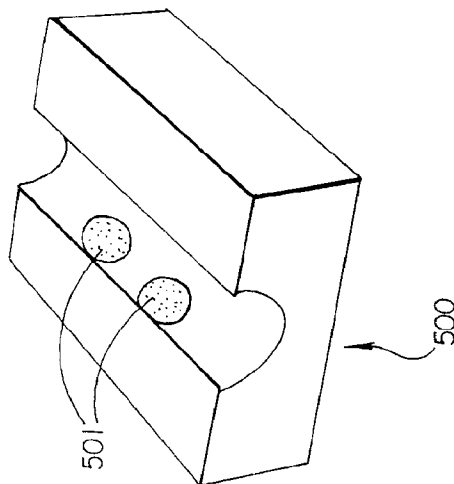


Fig. 13d

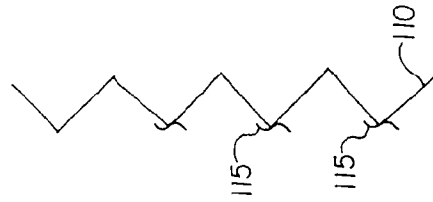


Fig. 13c

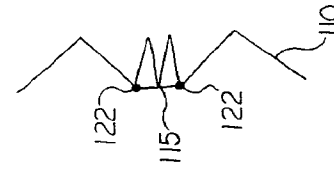


Fig. 13b

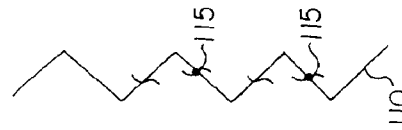


Fig. 13a



INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 00/00603

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61F2/06

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 98 11846 A (MICRO THERAPEUTICS INC) 26 March 1998 (1998-03-26) page 34, line 34 -page 37, line 27; figures 25-38	1
Y	US 5 192 307 A (WALL WILLIAM H) 9 March 1993 (1993-03-09) abstract; figures 7-12	1-12, 15
Y	US 5 779 732 A (AMUNDSON RODNEY R) 14 July 1998 (1998-07-14) column 5, line 21 - line 45; figure 2	1-12, 15
A	EP 0 382 014 A (ADVANCED CARDIOVASCULAR SYSTEM) 16 August 1990 (1990-08-16) abstract; figure 1A	1, 9
	-/--	

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

8 June 2000

Date of mailing of the international search report

16/06/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2

NL - 2280 HV Rijswijk

Tel. (+31-70) 340-2040, Tx. 31 651 epo.nl,

Fax: (+31-70) 340-3016

Authorized officer

Korth, C-F

INTERNATIONAL SEARCH REPORT

Inter national Application No

PCT/US 00/00603.

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	US 5 972 029 A (FUISZ RICHARD C) 26 October 1999 (1999-10-26) the whole document -----	1

INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/US 00/00603 .

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 9811846	A	26-03-1998	US 6007573 A AU 4484597 A	28-12-1999 14-04-1998
US 5192307	A	09-03-1993	US 5824038 A US 6015430 A US 5266073 A	20-10-1998 18-01-2000 30-11-1993
US 5779732	A	14-07-1998	NONE	
EP 0382014	A	16-08-1990	CH 678393 A CA 2008312 A DE 69014923 D DE 69014923 T JP 2931010 B JP 3198845 A US 5443500 A	13-09-1991 26-07-1990 26-01-1995 22-06-1995 09-08-1999 30-08-1991 22-08-1995
US 5972029	A	26-10-1999	NONE	